

WHAT IS CLAIMED IS:

Sub B7  
1. A method of forming a patterned thin film comprising:

5 depositing a thin film material on a surface of a substrate having thereon a patterned underlayer of a self-assembled monolayer.

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10 2. The method of claim 1, wherein said substrate is selected from the group consisting of: a metal, a metal oxide, a semiconductor, a metal alloy, a semiconductor alloy, a polymer, an organic solid, and a combination thereof.

15 3. The method of claim 2, wherein said substrate is an irregularly shaped substrate.

20 4. The method of claim 2, wherein said substrate is a solid substrate having a flexible, curved or planar geometry.

25 5. The method of claim 1, wherein said self-assembled monolayer is prepared by a process comprising the steps of:

providing a stamp having a surface;

Sub A1  
30 coating said surface of said stamp with an organic molecular species to produce a coated surface, said organic molecular species having a head functional group capable of interacting with said surface of said substrate, and a tail group for chemical

differentiation of said patterned and unpatterned regions of said coated surface;

placing said coated surface in contact with said substrate for a length of time sufficient to transfer  
 5 said self-assembled monolayer of said organic molecular species from said stamp to said substrate; and  
 removing said stamp.

6. The method of claim 5, wherein said stamp is  
 10 an elastomeric stamp.

7. The method of claim 5, wherein said stamp has  
 at least one indented and at least one non-indented  
 15 surface.

8. The method of claim 7, wherein said transfer  
 is in a pattern defined by the topography of said  
 stamp.

9. The method of claim 5, wherein said organic  
 molecular species has a functional head group selected  
 from the group consisting of: a phosphine, phosphonic  
 acid, carboxylic acid, thiol, epoxide, amine, imine,  
 hydroxamic acid, phosphine oxide, phosphite, phosphate,  
 20 phosphazine, azide, hydrazine, sulfonic acid, sulfide,  
 disulfide, aldehyde, ketone, silane, germane, arsine,  
 nitrile, isocyanide, isocyanate, thiocyanate,  
 isothiocyanate, amide, alcohol (hydroxyl), selenol  
 (selenide), nitro, boronic acid, ether, thioether,  
 25 carbamate, thiocarbamate, dithiocarbamate,  
 dithlocarboxylate, xanthate, thioxanthate,

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11. The method of claim 5, wherein said organic molecular species comprises one or more compounds selected from the group consisting of: a silane, a phosphonic acid, a carboxylic acid, a hydroxamic acid, a thiol, an amine, a phosphine, a hydrocarbon, partially halogenated hydrocarbon and a fully halogenated hydrocarbon.

12. The method of claim 5, wherein said organic molecular species comprises (tridecafluoro-1,1,2,2-tetrahydrooctyl)trichlorosilane.

13. The method of claim 5, wherein said organic molecular species comprises octadecylphosphonic acid.

14. The method of claim 1, wherein said self-  
5 assembled monolayer is prepared by a process comprising  
the steps of:

contacting said substrate and a solution comprising an organic molecular species having a head functional group capable of interacting with said surface of said substrate, and a tail group for chemical differentiation, said contacting being at a temperature and for a length of time sufficient to bind said functional head groups to said surface of said substrate; and

15 exposing said self-assembled molecular monolayer  
to radiation modulated spatially in intensity with a  
mask having one or more regions transparent to  
radiation to chemically modify said self-assembled  
molecular monolayer in a chemically distinct pattern  
20 defined by said transparent regions of said mask.

15. The method of claim 14, wherein said radiation is light.

25           16. The method of claim 14, wherein said mask is  
a photomask.

17. The method of claim 14, wherein said contacting is carried out by immersing said substrate in said solution comprising said organic molecular species.

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18. The method of claim 14, wherein said organic molecular species has a functional head group selected from the group consisting of: a phosphine, phosphonic acid, carboxylic acid, thiol, epoxide, amine, imine, hydroxamic acid, phosphine oxide, phosphite, phosphate, phosphazine, azide, hydrazine, sulfonic acid, sulfide, disulfide, aldehyde, ketone, silane, germane, arsine, nitrile, isocyanide, isocyanate, thiocyanate, isothiocyanate, amide, alcohol (hydroxyl), selenol (selenide), nitro, boronic acid, ether, thioether, carbamate, thiocarbamate, dithiocarbamate, dithlocarboxylate, xanthate, thioxanthate, alkylthiophosphate, dialkyldithiophosphate, and a combination thereof.

19. The method of claim 14, wherein said organic molecular species has a functional tail group selected from the group consisting of: a hydrocarbon, partially halogenated hydrocarbon, fully halogenated hydrocarbon, phosphine, phosphonic acid, carboxylic acid, thiol, epoxide, amine, imine, hydroxamic acid, phosphine oxide, phosphite, phosphate, phosphazine, azide, hydrazine, sulfonic acid, sulfide, disulfide, aldehyde, ketone, silane, germane, arsine, nitrile, isocyanide, isocyanate, thiocyanate, isothiocyanate, amide, alcohol (hydroxyl), selenol (selenide), nitro, boronic acid, ether, thioether, carbamate, thiocarbamate, dithiocarbamate, dithlocarboxylate, xanthate, thioxanthate, alkylthiophosphate, dialkyldithiophosphate, and a combination thereof.

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Figure 1. The effect of the concentration of the *Agrobacterium* suspension on the transformation efficiency of *Agrobacterium* strains. The *Agrobacterium* strains were transformed with the *Agrobacterium* suspension of 10<sup>6</sup> cells/ml (A), 10<sup>7</sup> cells/ml (B), 10<sup>8</sup> cells/ml (C), 10<sup>9</sup> cells/ml (D), 10<sup>10</sup> cells/ml (E), and 10<sup>11</sup> cells/ml (F). The transformation efficiency was determined by the number of transformants per 10<sup>6</sup> cells of the host strain. The data are the mean  $\pm$  SD of three independent experiments.

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26. The method of claim 23, wherein said long-chain organic polymer is polymethyl methacrylate/methyl methacrylate copolymer.

5 27. The method of claim 23, wherein said metallo-organic complex is tin 2-ethylhexanoate.

10 28. The method of claim 22, wherein said solution-based deposition process is a spin-coating process comprising the steps of:

flooding said substrate having thereon said patterned self-assembled molecular monolayer with a solution comprising a thin film material or a precursor thereof; and

15 spinning to deposit said thin film material thereby forming a patterned thin film on said substrate.

20 29. The method of claim 22, wherein said solution-based deposition process is an immersion-coating process comprising the steps of:

immersing said substrate having thereon said patterned self-assembled molecular monolayer into a solution comprising said thin film material, or a precursor thereof; and

25 withdrawing said substrate from said solution, thereby forming a patterned thin film on said substrate.

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Add A2

add B3